

EXHIBIT D

TWENTY-FOURTH EDITION

PHYSICIANS' DESK REFERENCE

to
PHARMACEUTICAL SPECIALTIES
and BIOLOGICALS

PATENT OFFICE

FEB 16 1970

SEARCH CENTER

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IN FIVE SECTIONS

An arbitrary page numbering plan is used to facilitate the compilation of this reference book.

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PUBLISHED BY MEDICAL ECONOMICS, INC.,
a subsidiary of

LITTON PUBLICATIONS, INC.
DIVISION OF LITTON INDUSTRIES
Oradell, New Jersey 07649



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Product Information

665

Pharmaceutical
(WENSWOOD AVE.
N.L. 50640)

In each tablet, active
ingredient is as follows:

1. Trigonitis, urethral
infections where
the bacteria are susceptible
to methylene blue, and
depends upon an
antibiotic as a prophylactic
agent in the treatment of
symptomatic infections
with specific therapy.

d relief of pain, relief
in of atropine and
content decreased.
on and Dosage: Adult,
four times per day
id intake. Acute
tablets every hour
by the recom-

half the adult dose.
Neither irritation
actions have been
pronounced dryness
ing, or difficulty
occurs, decrease
tixation or blurring
tinue use immediately
may be present
teropathy.

Administer with
known idiosyncrasy
patients suffering from
under this therapy
ions should be advised.

ions: Glaucoma,
k or pyloric ob-
struction and car-
ity to any of the in-
cool place. Owing
ture of the compo-
it is recommended
in glass.
Bottles of 100, 500

available: Yes.

dated Midland
orporation
N STREET
N.Y. 10509

ton)

Long-acting capsules
diphenhydramine
ve irritation caused
a sodium salt). Fe-
sion: The anticonvul-
le (almost free from
effects) usually lasts
he severity and fre-
quency. Improves
d patterns of epi-
lepsy, ability to concen-
trate.

and Dosage: One capsule,
the morning, is usually sufficient
the morning in most cases.
Doses higher than 500 mg.
may cause headache, drowsiness,
nausea and digestive and renal
disturbances, blood dyscrasias,
rash, jaundice, a lymphoma-like
hyperplasia, peripheral neuritis, arrhythmia,
and other conditions have been reported.
Gradual withdrawal is recom-

indications: Severe liver and renal
dysfunction and blood dyscrasia.
Bottles of 30, 100 and hospital
supply.

ACNAVEEN®
(Sodium Salicylate) acid (buffered)

Composition: Coated tablets with 0.75 Gm.
of sodium salicylate and 25 mg. INH.

Action and Uses: Antituberculous activity
against Mycobacterium tuberculosis and
delayed when INH and PAS
are used simultaneously. Tuberculous in-
fection should be treated with this product
in combination with streptomycin or
isoniazid.

Contraindications and Dosage: Recommended
dosage for adults: Three to four tablets, four
times daily, preferably before meals. This
dose provides 300 to 400 mg. INH and
0.75 Gm. buffered PAS.

Warnings: INH: CNS stimulation (can be
prevented with pyridoxine); peripheral neuritis
(can be prevented with pyridoxine);
allergic rash and fe-
brile reaction; hepatic depression,
hepatic disturbance, prolonged clotting
time, and psychotic reactions
have been reported.

Indications: Severe liver or renal
dysfunction, clinical and latent epilepsy, psy-
chotic and blood dyscrasias.
Bottles of 500 and 1000
coated tablets. Hospital packing.

ACNAVEEN'S SOLUTION® w/Vitamin A

Composition: A coating combination of
alcohol 2%, benzyl cinnamate 1.8%,
cinnamate 0.5% in olive oil with 10-
units of Vitamin A per cc.
Action and Uses: Eucrotic tissue formation
and necrotic resolution of local-
ized inflammation. Indicated in pelvic in-
flammatory disease, diabetic and athero-
sclerotic retinopathies and in deafness and
other inflammatory lesions of mid-
dle ear when inflammatory factors
are underlying factors.

Administration and Dosage: Intramuscular
injection only. See package circular for
dosage schedule.

How Supplied: Boxes of 12 and 50, 1 cc.

Cooper Laboratories, Inc.
Division of Aveeno Dermatologicals,
Wynne Pharmaceuticals, and
Wynne Laboratories, Inc. (Division)
1000 W. WENSWOOD AVENUE
N.L. 50635

ACNAVEEN® BAR
(Sodium Salicylate) acid (buffered)

Composition: ACNAVEEN BAR: Aveeno®
Colloidal Oatmeal (50%); a gentle, effec-
tive antipruritic; sulfur (2%); salicylic acid
(2%); hexachlorophene (2%). The pH of
the preparation has been adjusted to approximate
normal skin.

ACNAVEEN CREAM: Aveeno Colloidal
Oatmeal (10%); sulfur (2%); salicylic acid
(2%); hexachlorophene (1%) in a soap-
free base containing a blend of sudan-
ese and waxes.

Action and Uses: For gentle, safe, cleansing

in acne, oily skin and scalp. Acnaveen
therapy effectively cleanses the skin of
dirt, debris and excess sebum, while pro-
viding antipruritic and anti-irritative ac-
tions through the soothing effects of
Aveeno Colloidal Oatmeal. Especially use-
ful where the skin is irritated or sensitive.

Administration and Dosage: Acne—Wet
face thoroughly and massage into skin vig-
orously to produce lather. Allow to remain
on skin for several minutes. Rinse well. Re-
peat 2-3 times daily.

Shampoo—Wet hair thoroughly, then mas-
sage a liberal amount of Acnaveen Cream
into hair and scalp. Allow to remain five
minutes. Rinse. Repeat shampoo and rinse
immediately. Use twice weekly, or as re-
quired.

How Supplied: Acnaveen Bar—3.5 oz. Ac-
naveen Cream—4 oz. plastic tube.

AMINODUR® DURA-TAB® 4 1/4 gr. B
(Formerly Aminophylline Dura-Tab)

Composition: Each Aminodur Dura-Tab
contains aminophylline U.S.P. 4 1/4 gr. (ap-
prox. 0.3 Gm.) in a tablet matrix specially
designed for prolonged release of the drug.
Action and Uses: For use in the treatment
of chronic bronchial asthma and pulmo-
nary emphysema. Aminophylline has the
general physiological properties of the-
ophylline. Its chief use depends upon its en-
hancement of bronchial flow, relaxation of
bronchial and other smooth musculature,
and stimulation of the respiratory center.
Precautions: Use cautiously in patients
with poor renal function as a decreased
rate of excretion may lead to untoward
reactions.

Side Effects: Patients may develop gas-
trointestinal disturbances after prolonged
use. The simultaneous administration of
aluminum hydroxide decreases the incide-
nce of this side effect.

Administration and Dosage: Adults, 1 to 2
Aminodur Dura-Tabs every 8 to 12 hours
approximately 30 minutes before eating.
How Supplied: Aminodur Dura-Tabs,
scored, in bottles of 100 and 250.

AVEENO® BAR (Soap-free Cleansing Bar)

Composition: Aveeno® Colloidal Oatmeal
50%, anionic sulfonate, lanolin derivative,
hexachlorophene 2%.

Action and Uses: For use in housewives'
and infantile eczema, atopic and contact
dermatitis, dry skin, and prophylactically
in patients with sensitive skin, or wherever
soap or common detergents must be
avoided. Aveeno-Bar has a pH approx-
imating that of normal skin. It contains a
mild surfactant and hypoallergenic lanolin
derivative to provide gentle cleansing with
a soft and silky feel.
How Supplied: 3 1/2 oz. bar.

AVEENO® COLLOIDAL OATMEAL
(for soothing antipruritic colloidal baths)

Composition: Concentrated colloid produc-
ing fraction of oatmeal produced especially
for skin therapy.

Action and Uses: For use in acute, wet or
inflamed dermatoses, such as poison ivy,
poison oak, intertrigo, diaper rash and
atopic dermatitis. Particularly valuable
where soothing and anti-pruritic action is
desired.

Administration and Dosage: One cup to a tub
of warm water once or twice daily, or as
required. For infants, add 1-2 tablespoon-
fuls to bathwater.

How Supplied: 1 lb. and 4 lb. boxes.

F.S.N. 5505-200-8288

AVEENO® LOTION

Composition: Aveeno Colloidal Oatmeal
(10%) in a soothing aqueous lotion base,
containing propylene glycol, isopropyl my-
ristate and liquid petrolatum.

Action and Uses: A soothing, flexible, film-
forming shake lotion, providing symp-

tomatic relief of sunburn, poison ivy,
diaper rash, and other acute and subacute
dermatoses. It is compatible with many
agents including menthol, phenol, resorcinol,
sulfur, salicylic acid, liquor carbonis deter-
gens, hydrocortisone, etc. Avoids over-
drying. Cosmetic coloring blends with most
skin tones.

Administration and Dosage: Apply to af-
fected area 2 or 3 times daily, or as physi-
cian directs. Rub in gently.

How Supplied: 8 fl. oz. plastic bottles.

AVEENO® ORATED

Composition: Aveeno® Colloidal Oatmeal
impregnated with 35% emollient oils (liq-
uid petrolatum and hypoallergenic frac-
tions of lanolin).

Action and Uses: For use in chronic, dry
and subacute dermatoses where anti-
inflammatory and emollient effects are
desired, particularly itching dermatoses
such as senile pruritus, bath itch and
chicken pox.

Administration and Dosage: To a tub of
warm water, add 4 rounded tablespoonfuls
once or twice daily, or as required.

How Supplied: 10 oz. and 2 lb. boxes.

AVENOL™ BATH ADDITIVE

Composition: Liquid petrolatum, beeswax,
Aveeno® Colloidal Oatmeal, dewaxed
lanolin derivative, emulsifier.

Action and Uses: Combines the emolliency
of a bath oil with the soothing, relief-pro-
moting properties of Aveeno Colloidal
Oatmeal. For dry, itching skin conditions
such as senile pruritus, winter itch, atopic
dermatitis, psoriasis, ichthyosis, chicken
pox and nonspecific dry skin disorders.

Administration and Dosage: Add 2-3 cap-
sules to the bath. Disperse well before en-
tering tub. For infant or sponge bath use
1-2 capsules in bath water. Pat skin dry.

How Supplied: 8 fl. oz. plastic bottles.

ELIXOPHYLLINE®

Composition: Each 15 cc. (tablespoonful)
contains theophylline (anhydrous) 80 mg.,
alcohol 20% in a palatable, aromatic base.
Action and Uses: For symptomatic relief of
bronchial asthma, pulmonary emphysema
and other pulmonary diseases with bron-
chospasm, angina pectoris and cardiac
dyspnea.

Contraindications: May be contraindicated
in peptic ulcer.

Precautions: Do not use concurrently with
other theophylline preparations.

Note: Diabetic Information: Each 15 cc.
(tablespoonful) contains approximately 20
calories, 0.9 Gm. carbohydrate.

Administration and Dosage: SEVERE
ASTHMA ATTACK: Adults, 75 cc. (5 ta-
blespoonfuls). Children, 0.5 cc. per pound
of body weight. Do not repeat within six
hours. MAINTENANCE 24-HOUR
THERAPY: Adults, for first 6 doses—45 cc.
(3 tbsp.) before breakfast, at 3 p.m., and at
bedtime, then 30 cc. (2 tbsp.) doses at the
same times. Children, 0.5 cc. per pound of
body weight 3 times daily as above, then
0.2 cc.

How Supplied: Bottles of 16 fl. oz., 32 fl.
oz., 1 gal., a red, pleasant tasting solution.

ELIXOPHYLLINE-KI

Composition: Each 15 cc. (tablespoonful)
contains theophylline (anhydrous) 80 mg.,
potassium iodide 130 mg., alcohol 10%; in a
palatable, aromatic base.

Indications: For excessive tenacious mucus
in chronic asthma, severe, chronic and al-
lergic bronchitis, chronic obstructive pul-
monary emphysema.

Contraindications: Contraindicated in pa-
tients with hyperthyroidism or known sen-
sitivity to iodides. May be contraindicated
in peptic ulcer or gout.

Continued on next page

BBW 004405

666

Product Information

Always consult

Cooper—Cont.

Side Effects: Possible erythema, slight rhinitis, mild sore throat. If these symptoms develop, discontinue use.

Precautions: Do not use other theophylline preparations concurrently. Caution is recommended in patients during pregnancy. In some patients prolonged use of iodides can lead to hypothyroidism.

Note: Diabetic Information: Each 15 cc. (tablespoonful) contains approximately 24.5 calories, 4 Gm. carbohydrate.

Dosage: Adults, 30 cc. (2 tablespoonfuls) t.i.d. on arising, at 3 p.m. and on retiring. Children, 0.2 cc. per pound of body weight, t.i.d. as above.

How Supplied: Bottles of 8 fl. oz. and 32 fl. oz. (one quart)

EMULAVE™ BAR
(Cleansing Bar for Dry Skin)

Composition: A combination of vegetable oils and dewaxed lanolin fraction (25%), Aveeno® Colloidal Oatmeal (22%) and hexachlorophene (2%) in solid bar form containing a blend of soothing and wetting agents.

Action and Uses: A completely soapless cleansing bar for dry skin conditions, containing 25% emollients, which deposit a persistently adherent lipid moisture barrier on the skin. In addition, the Aveeno Colloidal Oatmeal content provides a gentle, soothing effect.

How Supplied: 3 oz. bar.

ENURETROL®

Composition: Each tablet contains: Ephedrine Sulfate—7.5 mg., Atropine Sulfate—0.15 mg., in a palatable, chewable base.

Action and Uses: As an adjunct for the control of nocturnal (functional) enuresis.

Effects: 1. Reduces bladder pressure, and also diminishes the amplitude and frequency of bladder contractions, thus increasing its capacity.

2. Stimulates the internal vesical sphincter, thus allowing the bladder to retain its contents.

Contraindications: Acute glaucoma, prostatic hypertrophy, or hypersensitivity to parasympathetic depressants or sympathomimetic agents.

Precautions: Should be used with caution in patients with coronary or cardiovascular disease and/or severe hypertension, hyperthyroidism, or diabetes mellitus.

Adverse Reactions: In large doses, effects such as rapid pulse, blurred vision, dryness of mouth, dizziness, flushing, constipation or sleeplessness may occur.

Administration and Dosage: Should be administered twice daily according to the age and weight of the patient for a minimum period of 4 to 6 weeks. A suggested initial dosage is: children 5 to 10 years, 1 tablet in the morning and 1 at bedtime; children 11 to 15 years, 2 tablets in the morning and 2 at bedtime; patients over 15 years, 3 tablets in the morning and 3 at bedtime. Once the patient no longer exhibits symptoms of enuresis, gradually curtail medication over a two-week period. At the end of the initial 4 to 6 week period, if the patient has not responded to therapy under careful supervision of the physician, dosage can be raised to the limits of tolerance. (Tolerance limits are readily recognized by the appearance of blurred vision, dryness of mouth, flushing, etc.) Best results with Enuretrol are obtained if bladder training is practiced on a regular basis by the patient during the daytime medication period. The patient should drink larger than normal amounts of liquids during the day and retain urine to the point of discomfort. This will cause the bladder to distend and thereby increase its capacity. However, as dry a diet as possible

should be used during the evening hours. Tablets may be swallowed, chewed or, where indicated, crushed in a little water.

How Supplied: Enuretrol is supplied in bottles of 50 and 500 scored, cherry-flavored, chewable tablets.

ERGOMAR®
(ergotamine tartrate)

Composition: Each sublingual tablet contains specially processed ergotamine tartrate, 2.0 mg.

Indication: Migraine.

Effects: Ergomar (ergotamine tartrate) exerts a direct effect on cranial blood vessels, causing vasoconstriction with concomitant decrease in the pulsations probably responsible for migraine symptoms. It is thus generally considered to be a specific agent for the therapy of this condition.

Administration and Dosage: All efforts should be made to initiate therapy as soon as possible after the first symptoms of the attack are noted, since success is proportional to rapidity of treatment, and lower dosages will be effective. At the first sign of an attack, or to relieve the symptoms of the full-blown attack, one tablet is placed beneath the tongue. Another tablet should be taken at half-hourly intervals thereafter, if necessary, but dosage must not exceed 3 tablets in any 24-hour period. Limit dosage to not more than 10 mg. in any one week.

Side Effects: No serious complications have been reported from the use of Ergomar (ergotamine tartrate) in the absence of contraindications and in recommended dosages. Unpleasant side effects which may occur include nausea and vomiting, weakness in the legs, muscle pains in the extremities, numbness and tingling of fingers and toes, precordial distress and pain, and transient tachycardia or bradycardia. Localized edema and itching may occur in the rate sensitive patient. Side effects are usually not such as to necessitate interruption of therapy.

Precautions and Contraindications: Avoid prolonged administration or dosage in excess recommended because of the danger of ergotism and gangrene. Contraindicated in sepsis, occlusive vascular disease (thromboangiitis obliterans, luetic arteritis, severe arteriosclerosis, coronary artery disease, thrombophlebitis, Raynaud's disease), hepatic disease, renal disease, severe pruritus, and pregnancy.

FERRONORD®

(brand of ferroglycine sulfate complex)*

Composition: Each enteric coated tablet contains: 250 mg. of ferrous iron amino acid complex equivalent to 40 mg. Ferrous Iron.

Action and Uses: For treatment of iron deficiency anemias.

Adult Dosage: One tablet 3 to 4 times daily, after meals and before retiring.

Children's Dosage: 3 to 12 years; one tablet 2 to 3 times daily after meals. For higher dosage or for younger children, as directed by physician.

How Supplied: Bottles of 100's.

*U.S. Patent Nos. 2,877,253; 2,957,806

FERRONORD®-DLA

(brand of ferroglycine sulfate complex)*

Composition: Each capsule contains 400 mg. of ferrous iron-amino acid complex (equivalent to 75 mg. Ferrous Iron).

Action and Uses: For treatment of iron deficiency anemias.

Dosage: 1 or 2 capsules daily or as directed by a physician.

How Supplied: Bottles of 30's.

*Patent Nos. 2,877,253; 2,957,806.

KAY CIEL™ Elixir

(potassium chloride)

Composition: Each 15 cc. (one tablespoonful) contains potassium chloride 1.5 Gm.

supplying 20 mEq. of elemental potassium in a cherry-flavored, palatable, base alcohol 4%. Contains no sugar.

Action and Uses: For the treatment of potassium deficiency occurring during thiazide diuretic or corticosteroid therapy, digitalis intoxication, low intake of potassium, or as a result of severe vomiting and diarrhea.

Contraindications: Impaired renal function, untreated Addison's Disease, dehydration, heat cramps, and hyperkalemia.

Precautions: Potassium chloride should be administered with caution and adjust the requirements of the individual since the amount of deficiency and responding daily dose is often not known. Excessive or even therapeutic doses may result in potassium intoxication. The patient should be checked frequently, periodic ECG and/or plasma potassium levels made. High plasma concentrations of potassium ion may cause cardiac arrest, arrhythmias or arrest. Use with caution in patients with cardiac disease, hypokalemic states, attention should be directed toward the correction of the frequently associated hypochloremia.

Adverse Reactions: Vomiting, nausea, abdominal discomfort, and diarrhea may occur.

Administration and Dosage: Adults: One tablespoonful (15 cc.) diluted in one of water, tomato or orange juice, daily after morning and evening meals. Larger doses may be indicated according to the individual patient's requirements should be administered under close supervision, due to the possibility of potassium intoxication. Patients should be cautioned to follow directions explicitly in regard to dilution of KAY CIEL Elixir to avoid gastrointestinal injury.

Toxicity: Symptoms and signs of potassium intoxication include muscle weakness, confusion, paresthesia of the extremities, weakness of the legs, flaccid paralysis, in blood pressure, cardiac arrhythmias, heart block. When hyperkalemia exists should be promptly treated with the continuance of potassium administration, other steps to lower serum levels indicated, since sudden shift in plasma may induce potentially dangerous arrhythmias.

How Supplied: 4 fl. oz., 16 fl. oz. (one and 128 fl. oz. (one gallon) bottles.

LUASMIN® CAPSULES

Description: Each blue and white capsule contains phenobarbital, sodium (which may be habit-forming) 30 mg. (½ theophylline sodium acetate 0.3 Gm. gr.) equivalent to 0.12 Gm. theophylline sodium acetate 30 mg. (½ gr.).

Action and Uses: For prophylaxis and temporary relief of bronchial spasms, unique buffered xanthine derivative theophylline sodium acetate, together with phenobarbital provides bronchodilation and fast relief of bronchospasm. The full therapeutic dose of phenobarbital, helps allay anxiety and mood to some extent, the possible stimulant effects of the bronchodilating agent.

Contraindications: Porphyria and sensitivity to any of the ingredients.

Precautions: Use cautiously in patients with cardiovascular disease, severe hypertension, hyperthyroidism, prostatic hypertrophy, or glaucoma. In combination with digitalis, serious disturbances of rhythm may occur. As with all medications, use cautiously in pregnant patients, especially during the first trimester.

Adverse Reactions: Mild gastrointestinal irritation, difficulty in voiding, palpitations, insomnia, nervousness and drowsiness occasionally occur.

Dosage and Administration: Adults: One capsule every 3 to 4 hours.